

Mitochondrial Mechanisms Of Degeneration And Repair In Parkinsons Disease

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Mechanisms of mitochondrial dysfunction and their impact on age-related macular degeneration. 1. Introduction to RPE degeneration and development of AMD. AMD is associated with several environmental and genetic risk factors that are linked to ... 2. RPE mitochondria damage and dysfunction during ...

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Mechanisms of mitochondrial dysfunction and their impact ...

In fact, the most common approach to the generation of animal and cell-culture models of idiopathic Parkinson's disease involves exposure to mitochondrial toxins. Even more compelling is the fact that most familial patients harbor genetic mutations that cause disruptions in normal mitochondrial morphology and function.

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Oxidative stress-induced damage to the retinal pigment epithelium (RPE) is considered to be a key factor in age-related macular degeneration (AMD) pathology. RPE cells are constantly exposed to oxidative stress that may lead to the accumulation of damaged cellular proteins, lipids, nucleic acids, and cellular organelles, including mitochondria.

Mechanisms of mitochondrial dysfunction and their impact ...

Similarly, loss of the AAA+ protease YME1L, which not only degrades damaged or non-assembled proteins in the mitochondrial inner membrane but also cleaves Opa1 to activate it and catalyze inner membrane fusion, causes ocular degeneration and axonal degeneration in the spinal cord. The potential role of alterations in function of AAA+ proteases in muscle and motoneuron mitochondrial impairment with aging is unknown, but is worthy of consideration based upon the aforementioned effects.

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Manifestations of mitochondrial dysfunction with aging include impaired respiratory function, elevated reactive oxygen species production, and increased susceptibility to permeability

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In this respect, mitochondria are implicated in aging and age-related neurodegenerative disorders, and are also likely key to aging muscle changes through their direct effects in muscle fibers and through secondary effects mediated by mitochondrial impairments in motoneurons.

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The main pathways leading to mitochondria-associated cellular dysfunction include (1) calcium overload in the matrix and mPTP pore opening, (2) cytochrome c release and activation of apoptosis ...

Mitochondrial dysfunction in neurological disorders ...

Comment on: "Mitochondrial Mechanisms of Neuromuscular Junction Degeneration with Aging. Cells 2020, 9, 197" Cells. 2020 Jul 29;9(8):1796. doi: 10.3390/cells9081796. Authors Allen Herbst 1 ...

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Mitochondrial Mechanisms of Degeneration and Repair in Parkinson's Disease. Editors: Buhlman, Lori M. (Ed.) Free Preview. Summarizes theories of how poor mitochondrial function and morphology contribute to neurodegeneration; Discusses both sporadic and familial forms of Parkinson's disease ...

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A newly discovered self-destructive mechanism in mitochondria, the cells' powerhouses, may be one of the first deficits leading to motor neuron degeneration associated with toxic TDP-43 clumps — a hallmark of amyotrophic lateral sclerosis (ALS).

Mitochondrial 'Suicide' May Trigger Neurodegeneration in

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Degeneration of chronically demyelinated axons is a major cause of irreversible neurological disability in multiple sclerosis (MS) patients. Development of neuroprotective therapies will require elucidation of the molecular mechanisms by which neurons and axons degenerate. Methods

Mitochondrial dysfunction as a cause of axonal ...

Author summary Phosphatidylethanolamine (PE) is a critical component of all cellular membranes, and maintaining cellular PE homeostasis is critical for survival and function of cells especially neuronal cells. There are two major PE synthesis pathways in eukaryotes, the CDP-ethanolamine pathway in the endoplasmic reticulum (ER) and the PSD pathway in mitochondria.

PE homeostasis rebalanced through mitochondria-ER lipid

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In this respect, mitochondria are implicated in aging and age-related neurodegenerative disorders, and are also likely key to aging muscle changes through their direct effects in muscle fibers and through secondary effects mediated by mitochondrial impairments in motoneurons.

Mitochondrial Mechanisms of Neuromuscular Junction ...

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potential is required for ER fragmentation and mitochondrial Ca^{2+} elevation during neuritic degeneration. Mechanically, tightening of the ER-mitochondria associations by expression of a short "synthetic linker" and ER Ca^{2+} releasing together could promote mitochondrial.

Mitochondrial Membrane Potential-dependent Endoplasmic ...

The specific degeneration of neurons in the substantia nigra pars compacta causes rigidity, tremor, and bradykinesia, all of which are hallmark symptoms of Parkinson's disease. One of the potential molecular factors that might induce the loss of dopaminergic neurons is an elevated level of reactive oxygen species which causes cell damage over ...

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